

Novel immunity proteins associated with colicin M-like bacteriocins exhibit promiscuous protection in *Pseudomonas*

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Novel immunity proteins associated with colicin M-like bacteriocins exhibit promiscuous protection in *Pseudomonas*

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Provisional

Abstract

Bacteriocins related to colicin M, acting via cleavage of the cell wall precursor lipid II, have been characterized in γ - and β -proteobacteria. Depending on the species, immunity is provided by either an inner membrane-anchored periplasmic protein or by an integral membrane protein. In *Pseudomonas* however, the immunity partner of colicin M-like bacteriocins remains unknown. Based on an in silico analysis in pseudomonads genomes, we here identify a gene encoding a putative immunity partner that represents a novel type of integral membrane protein (PmiA, *Pseudomonas* colicin M-like immunity type A). By heterologous expression of *pmiA* genes in susceptible strains, we show that immunity to colicin M-like bacteriocins is indeed provided by the cognate PmiA. Sequence homology among PmiA proteins is essentially absent, except for a short motif with a conserved periplasm-exposed aspartate residue. However, PmiA's protective function is not abolished by changing this acidic residue to the uncharged alanine. Immunity by PmiAs appears promiscuous to the extent that PmiA homologues from a clade sharing less than 40% pairwise amino acid identity, equally provide protection against the bacteriocin linked to the original PmiA. This study shows that multiple immunity factors have evolved independently to silence lipid II-targeting enzymatic bacteriocins. Their relaxed bacteriocin immunization capacity contrasts to the strict specificity of immunity proteins shielding the enzymatic domain of nuclease bacteriocins. The nature of associated immune functions needs consideration when using such natural protein antibiotics or designing novel variants.

1 Introduction

Antibacterial proteins are currently subject of renewed interest in the search for antimicrobials with novel modes of action. As yet poorly explored potential sources of such alternative protein-based antibiotics include lytic bacteriophages (Gerstmans *et al.*, 2016) and bacteriocinogenic strains (Schulz *et al.*, 2015). Bacteriocins are key mediators of intra-species competition between bacteria and combine high specificity with single-hit killing potency. Prominent bacteriocin producers that target human and plant pathogenic Gram-negative bacteria are found among enterobacteria (Cascales *et al.*, 2007; Grinter *et al.*, 2014; Holt *et al.*, 2013; Schulz *et al.*, 2015) and pseudomonads (Ghequire & De Mot, 2014).

Pseudomonas bacteriocins are very diverse with respect to structure and mode of action. R-type and F-type tailocins are multiprotein complexes that are evolutionary related to bacteriophage tails (Ghequire & De Mot, 2015a; Ghequire *et al.*, 2015; Hockett *et al.*, 2015). Lectin-like bacteriocins are constituted of a tandem of monoco mannose-binding lectin domains, and kill target cells via an unknown mechanism (Ghequire *et al.*, 2013; Ghequire *et al.*, 2014; McCaughey *et al.* 2014). S-type bacteriocins are multi-domain toxins with a modular structure similar to colicins, the bacteriocins from *Escherichia coli* (Cascales *et al.*, 2007; Dingemans *et al.*, 2016; Ghequire & De Mot, 2014; McCaughey *et al.*, 2016). Self-inhibition of *Pseudomonas aeruginosa* strains producing a S-type bacteriocin (S pyocin) is circumvented by an immunity protein that transiently impedes the bacteriocin's toxic function. This immunity complement is co-expressed with the bacteriocin, and is typically encoded downstream of the pyocin gene, on the same or opposite strand. In the case of S-type bacteriocins with nuclease activity, bacteriocins are secreted as high-affinity bacteriocin-immunity protein complexes (Joshi *et al.*, 2015). *Pseudomonas* colicin M-like bacteriocins (PseuMs) host a lipid II-cleaving domain, homologous to the catalytic domain of colicin M (ColM) (Barreteau

et al., 2009; Barreteau *et al.*, 2012; Grinter *et al.*, 2012b). However, in contrast to S pyocin-immunity complexes, it is not clear how PseuM producers protect themselves. Inspection of their *pseuM* genomic context did not reveal homologues of *cmi*, the cognate immunity partner for ColM in *Escherichia coli*, nor were such homologues detected elsewhere in their genomes (Barreteau *et al.* 2009).

In *E. coli*, the *cmi* gene is located downstream of the bacteriocin gene on the opposite strand (Olschl ger *et al.*, 1984; Olschl ger & Braun, 1987). Located in the periplasm, Cmi is anchored to the inner membrane by an amino-terminal hydrophobic α -helix, assisting in secretion but not being cleaved off. There, Cmi transiently impedes colicin M's activity during secretion and inactivates imported ColM (Gross & Braun, 1996; Olschl ger & Braun, 1987; Olschl ger *et al.*, 1991). The crystal structure of Cmi revealed the presence of four α -helices and four β -strands, and is exemplary of a YebF fold (G rard *et al.*, 2011, Us n *et al.* 2012). In *Pectobacterium* a *cmi*-like gene is located downstream and opposite of the pectocin M gene, but its protective functionality remains to be demonstrated (Grinter *et al.*, 2012a). In most strains of *Burkholderia* producing colicin M-like burkhocins, immunity is mediated by *bmiB* genes that give rise to proteins with a small size and predicted topology similar to Cmi (Ghequire & De Mot, 2015b). However, the amino acid sequence of the periplasmic moiety of BmiB is unrelated to Cmi and lacks the characteristic YebF domain. Fewer strains carry a second type of burkhocin immunity gene (*bmiA*) that encodes a small integral membrane protein comprising three transmembrane helices.

In this study we investigated the genomic context of *pseuM* genes, in search for candidate immunity partners. We demonstrate that a gene downstream of *pseuM* provides cognate immunity. Apart from sharing a common membrane topology reminiscent of BmiA proteins, the encoded proteins display very poor sequence conservation across PseuM-bacteriocinogenic pseudomonads. The possible role of a short semi-conserved motif in these proteins is explored, as well as the capability of PmiA proteins to immunize targeted cells in an (a)specific way.

2 Materials and methods

2.1 Genome searches and phylogenetic analysis

PseuM bacteriocin genes in *Pseudomonas* genomes were identified by homology searches using the National Center for Biotechnology (NCBI) non-redundant database. The Pfam Colicin_M module PF14859 was used as a query. Genes downstream of unique *pseuM* bacteriocin gene products were selected for further analysis. Multiple sequence alignments were generated with MUSCLE and phylogenetic reconstructions were carried out with PhyML (1000 bootstrap replicates), implemented in Geneious v7.1.7. The JTT substitution model (Jones *et al.*, 1992) was used for studying the evolution of a large number of amino acid sequences. Amino-terminal signal sequences were predicted by PRED-TAT (<http://www.compgen.org/tools/PRED-TAT>) and TOPCONS (<http://topcons.cbr.su.se/>). Transmembrane regions and topology were predicted by TMHMM (<http://www.cbs.dtu.dk/services/TMHMM/>), and TOPCONS.

2.2 Bacterial strains and media

Strains used in this study are listed in Table S1. *Escherichia coli* was grown in LB (MP Biomedicals) at 37 C, whereas *Pseudomonas* strains were grown with shaking at 200 rpm in Trypticase Soy Broth (TSB, BD Biosciences) and Casamino Acids Medium (CAA, BD Biosciences) at 30 C, except for *P.*

97 *aeruginosa* that was grown at 37°C. Growth media were supplemented with agar (1.5%, Invitrogen),
 98 filter-sterilized isopropyl-β-D-thiogalactopyranoside (IPTG, 20 µg/ml, ForMedium), kanamycin (50
 99 µg/ml, Sigma-Aldrich), or tetracycline (15-150 µg/ml, Sigma-Aldrich) when required. Strains were
 100 kept on plate at 4°C, or at -80°C in glycerol (25% v/v, Sigma-Aldrich).

101 2.3 DNA methods and construction of plasmids

102 Genomic DNA was collected with the Puregene Yeast/Bact. Kit B (Qiagen). Synthetic genes and
 103 primers were obtained from IDT DNA. Genes encoding bacteriocins (ERS445055_00256 from *P.*
 104 *aeruginosa* NCTC10332, PFLQ8_1129 from *P. fluorescens* Q8r1-96 (Landa *et al.*, 2003) and
 105 PSPTO_0572 from *P. syringae* pv. *tomato* DC3000) and putative immunity proteins, encoded
 106 downstream, were amplified by polymerase chain reaction (PCR) with Q5 polymerase (New England
 107 Biolabs), with a T100TM Thermal cycler (Bio-Rad). Q5 polymerase was used according to the
 108 supplier's specifications. All primers are listed in Table S2. PCR fragments were purified with the
 109 Genelute PCR clean-up kit (Sigma-Aldrich), and double digested with NcoI/XhoI (1 hour at 37°C,
 110 New England Biolabs) in the case of the bacteriocin genes, and with PstI/XbaI (1 hour at 37°C, New
 111 England Biolabs) in the case of the putative immunity genes. The immunity gene amplicon of *P.*
 112 *aeruginosa* NCTC10332 was digested with PstI/EcoRI (1 hour at 37°C, New England Biolabs).
 113 Purified bacteriocin genes were ligated with T4 DNA ligase (1 hour at 37°C, Invitrogen) in pET28a
 114 (+) and putative immunity genes in shuttle vector pJB3Tc20. Ligation products were transformed to
 115 *E. coli* DH5α via heat shock. Standard methods were used for the preparation of competent *E. coli*
 116 cells and heat shock transformation of *E. coli* (Green and Sambrook, 2012). Transformants were
 117 initially verified for the presence of insert by PCR with *Taq* polymerase (New England Biolabs),
 118 using conditions as specified by the supplier. Plasmids were subsequently harvested with the
 119 Genelute HP plasmid miniprep kit (Sigma-Aldrich) and sequence confirmed for the presence of
 120 correct insert (GATC Biotech, Constance, Germany). Cloned immunity genes encompassed a 19-bp
 121 upstream region containing the predicted ribosome binding sites. Resulting plasmids are summarized
 122 in Table S1. The pET28a constructs were transformed to *E. coli* BL21/DE3 via heat shock, and
 123 pJB3Tc20 constructs electroporated to PseuM-susceptible *Pseudomonas* strains, and selected on the
 124 proper medium. Immunity genes encoding a mutation in a conserved aspartate residue were
 125 constructed via splicing by overlap extension, using pCMPG6251 (encoding the putative immunity
 126 complement of PseuM_{DC3000}), pCMPG6252 (complement of PseuM_{NCTC10332}), pCMPG6269
 127 (complement of PseuM_{Q8r1-96}) as a template.

128 2.4 Expression and purification of recombinant bacteriocins

129 Recombinant carboxy-terminally His-tagged bacteriocins were generated in *E. coli* BL21(DE3)
 130 carrying pCMPG6248 (encoding PseuM from *P. syringae* DC3000), pCMPG6250 (encoding PseuM
 131 from *P. aeruginosa* NCTC10332) or pCMPG6271 (encoding PseuM from *P. fluorescens* Q8r1-96).
 132 5-ml overnight cultures were transferred to 500 ml LB Erlenmeyer flasks and incubated at 37°C until
 133 OD₆₀₀ reached 0.7. After, cultures were cooled and supplemented with IPTG (1 mM final
 134 concentration) and incubated at 20°C for 16 hours. Next day, cells were harvested via centrifugation
 135 (20 min, 5000 g; Beckman X-15R) and frozen overnight (-20°C). Subsequently, the cell pellets were
 136 thawed, resuspended in lysis buffer (5 ml/g cell pellet; 300 mM NaCl, 50 mM NaH₂PO₄, 10 mM
 137 imidazole, pH 8.0) and sonicated (amplitude 20%, 10 cycles of 30 sec on/off; Branson Digital
 138 Sonifier). Samples were treated with nuclease (0.01 U/µl, 37°C, 1 hour; Invitrogen), remaining debris
 139 and insoluble proteins removed via centrifugation (30 min, 10000 g), and supernatants filtered (0.20
 140 µm, Sarstedt). Soluble fractions containing the recombinant proteins were loaded on a 5-ml HisTrap
 141 HP column (GE Healthcare) and purified by nickel affinity chromatography with an Äkta Purifier

(GE Healthcare). Matrix-bound proteins were eluted with a linear gradient of imidazole (10 mM-500 mM) in lysis buffer. Fractions eluted at high imidazole concentrations were validated for the presence of recombinant protein, pooled and dialyzed to Tris buffer (50 mM, NaCl 20 mM, pH 7.5). Concentration of purified PseuMs was determined with a Genesys 10S UV-Vis Spectrophotometer (Thermo Scientific). Calculated extinction coefficients were 30612 mol/L⁻¹ cm⁻¹, 33062 mol/L⁻¹ cm⁻¹ and 30367 mol/L⁻¹ cm⁻¹ for the His-tagged PseuM bacteriocins from strains DC3000, NCTC10332 and Q8r1-96, respectively.

2.5 Bacteriocin assay

Bacteriocin activity was determined via spot assay. Filter-sterilized 20-μl volumes of purified recombinant proteins (1 mg/ml) were spotted onto bacterial cell lawns. After, samples were air-dried and incubated overnight. Dialysis buffer was used as a negative control. Following day, petri dishes were scored for zones lacking bacterial growth (halos).

2.6 Nucleotide Accession Number

GenBank accession number of the nucleotide sequence of the *pseuM-pmiA* pair from *P. putida* RW10S2 (Rokni-Zadeh *et al.*, 2012) is KX086738.

3 Results

3.1 *In silico* search for candidate genes providing immunity to *Pseudomonas* colicin M-like bacteriocins

Based on the genetic architecture of modular bacteriocins in Gram-negatives, it is anticipated that bacteriocin immunity genes are likely to be found in proximity of their corresponding toxin genes. By scrutiny of their genomic contexts we observed that the characterized *Pseudomonas* colicin M-like bacteriocin genes, as well as several other predicted PseuM-encoding genes (Figure S1A), are consistently followed on the same DNA strand – at variable distances but within < 200 nucleotides – by an open reading frame, but infrequently annotated as a coding region (Table S3). Typically, G+C content of these regions (~1.8 kb) is significantly lower (41-52%) than the *Pseudomonas* genome averages (*P. aeruginosa*: 66.2%, 1647 genomes; *P. fluorescens*: 60.2%, 92 genomes; *P. syringae*: 58.7%, 153 genomes). None of the deduced protein sequences contains the YebF domain signature (Pfam PF13995) as found in Cmi and the putative pectocin M immunity proteins.

In fourteen *Pseudomonas* strains, mostly *P. syringae* pathovars, the encoded proteins (encompassing eight unique but very similar sequences of ~100 AAs) show only borderline homology with the burkhocin-associated BmiB immunity proteins (12-24% pairwise AA identity). However, these pseudomonad proteins, designated PmiBs, share with BmiBs two perfectly conserved cysteines (Figure S2). The PmiB and BmiB proteins are predicted to be translocated to the periplasm, albeit this appears to be achieved in different ways: instead of the BmiB amino-terminal membrane-anchoring segment or Sec-dependent cleavable signal sequence, a cleavable lipoprotein signal sequence typifies PmiB. This common feature is reflected in the phylogeny of ColM domains that reveals a well-separated clade harboring both the burkhocins and this small subset of PseuM proteins (Figure S1B).

The majority of PseuM-derived ColM domains constitutes a separate clade of much more diversified sequences that are evolutionarily related to the corresponding domains in colicin M and colicin M domain-containing pectocins (Figure 1A, Figure S1B). In this second subset of PseuM-bacteriocinogenic strains the bacteriocin gene is linked with a gene encoding a protein unrelated to PmiB (~137 AAs; Figure S3). This protein family, designated PmiA, features four transmembrane helices, of which the first is predicted to represent a candidate Sec- or Tat-dependent cleavable signal sequence (Figure 2A). The inferred topology, with two moieties facing the periplasm and two stretches (including the carboxy-terminus) residing in the cytoplasm, is reminiscent of the one adopted by BmiA proteins, albeit the latter lack an equivalent secretory signal sequence (Ghequire & De Mot, 2015b). No sequence homology can be detected between BmiAs and PmiAs however. Notably, four phytopathogenic strains of the *P. syringae* group each carry two unlinked PseuM-encoding paralogues, one from each PseuM clade and clustered with, respectively, a *pmiA* and *pmiB* gene. The low level of sequence identity between these bacteriocins (~24%) combined with the different nature of the respective candidate immunity proteins argues against a common ancestry and favors their independent acquisition.

In different strains, the *pseuM-pmiA* gene pair has been recruited to diverse genomic locations. Several of these regions correspond to mobile elements. In *P. aeruginosa*, the gene couple generally occurs on ExoU island A, as has previously been noted for the *pseuM* of *P. aeruginosa* JJ692 (Barreteau *et al.*, 2009). In some strains from the *P. syringae* group, such as isolates 642 and LMG 2367, *pseuM-pmiA* combinations appear as a cargo gene couple in a *trpE/trpG*-integrated Rp4 tailocin cluster (Hockett *et al.*, 2015). In other *P. syringae* pathovar strains, such as DC3000 and ICMP 3923, the cargo is equally part of a Rp4-type sequence, although incomplete and apparently linked with diverged sequences. In some *P. fluorescens* group strains, such as *P. brassicacearum* NFM421 and 51MFVCI2.1, and *P. fluorescens* Q8r1-96, the gene pair has been recruited by a Rp4 tailocin cluster as well, sandwiched between *mutS* and *cinA* (Ghequire *et al.*, 2015; Hockett *et al.*, 2015). For other strains, such as *P. veronii* R4, *Pseudomonas* sp. BRG-100 and *P. tolaasii* 6264, regions flanking the *pseuM-pmiA* couples correspond to an unassigned prophage. The presence of bacteriocin cargo genes on mobile elements has previously been noted for lectin-like bacteriocins embedded in *Pseudomonas* Rp2 tailocin clusters (Ghequire *et al.*, 2015).

Some *Pseudomonas* strains carry a *pmiA* gene that is not linked to a cognate *pseuM* gene. The presence of *pseuM* remnant sequences upstream of such a *pmiA* orphan indicates that these strains may represent so-called cheaters that have lost the capacity of bacteriocin production but retain the associated immune function (Figure 3). For instance, in the type strain *Pseudomonas lundensis* (DSM 6252; De Jonghe *et al.* 2011) a frameshift impairs functionality of a *pseuM* gene that has remained intact in *Pseudomonas* sp. AU9518. From the equivalent genomic region of type strain *Pseudomonas weihenstephanensis* (DSM 291166; von Neubeck *et al.* 2016) and some related strains (TAD18; TAA207 and DSM 28140, not included in Figure 3), most of the 3' region of a *pseuM* gene has been deleted, while their *pmiA* genes are well conserved (>92% AA identity). No obvious *pseuM* sequences remain adjacent to the *pmiA* gene of Arabidopsis isolate *Pseudomonas* sp. Leaf48 (Bai *et al.* 2015), that is flanked by prophage genes.

3.2 PmiA proteins are cognate PseuM immunity partners

Previously described PseuM bacteriocins from *Pseudomonas aeruginosa* NCTC10332 (PaeM), *Pseudomonas fluorescens* Q8r1-96 (PflM) and *Pseudomonas syringae* pv. *tomato* DC3000 (PsyM), each with a downstream *pmiA* gene (as identified by *in silico* analysis described in section 3.1), were recombinantly expressed in *E. coli* and the His-tagged proteins purified by Ni-NTA affinity

chromatography. Strain NCTC10332 encodes a PseuM bacteriocin that shares 90% AA identity with PaeM from *P. aeruginosa* JJ692 (Barreteau *et al.*, 2009). Antibacterial activity of the recombinant proteins was challenged via spot-on-lawn assay against a panel of *Pseudomonas* strains, and strains with a clear halo susceptibility phenotype were selected for further experiments. Such indicator strains identified for PaeM, PflM and PsyM were *P. aeruginosa* CPHL12447, *P. fluorescens* F113 and *P. syringae* pv. *lachrymans* LMG 5456 (Grinter *et al.*, 2012b), respectively.

Next, the putative immunity genes from strains NCTC10332, Q8r1-96 and DC3000 were cloned in shuttle vector pJB3Tc20, and introduced in the corresponding PseuM-susceptible strains. Transformants were tested for altered PseuM sensitivity, using strains carrying empty vectors as negative controls (Table 1). When equipped with the cognate *pmiA*, transformants became fully insensitive to the respective bacteriocin (tested at 1 mg/ml by spot assay). When indicator strains were provided with either of the other two immunity genes, no diminished PseuM inhibition was observed. Together this indicates that PmiAs are cognate immunity partners of the bacteriocins they are associated with. Purified membrane fragments with the gene product of *pmiA*_{DC3000} were analyzed via Maldi-MSMS (Poetsch *et al.*, 2008), but no peptide fragments assignable to the PmiA of interest could be retrieved (data not shown).

Since colicin M and colicin M-like bacteriocins domains exert their action in the periplasm, it is plausible that (at least part of) the periplasm-exposed moieties of PmiA proteins participate in the immunity function, directly or indirectly impeding the catalytic function of the ColM domain. Conserved residues in the immunity module may be critical in providing such property. In the mature PmiA proteins, a short well-conserved stretch with consensus motif D[T/S]XGXP precedes the second predicted transmembrane segment (Figure 4, Figure S3). No obvious sequence conservation among PmiA proteins can be detected for the shorter second periplasm-exposed loop. To further scrutinize the possible role of the first periplasmic stretch, the conserved Asp residue was mutated to Ala in the PmiAs of strains NCTC10332, Q8r1-96 and DC3000. Upon introduction of the PmiA variants in the respective indicator strains, the immunity phenotype to the respective PseuMs was evaluated. Halos were absent for all three indicator strains after bacteriocin spotting, showing the same phenotype as caused by the native immunity gene products (Table 1). This suggests that the conserved Asp residue is not instrumental to PmiA's immunity function.

3.3 Immune promiscuity among sequence-diverged PmiAs

Noteworthy, sequence divergence of the PseuM-PmiA toxin-immunity pairs seems to be most pronounced among strains belonging to the *P. fluorescens* group. In contrast, for *P. syringae* pathovar isolates, high conservation of the ColM domain (at least 94% pairwise AA identity) (Figure S1A, Figure S1B) is reflected in well conserved immunity proteins (at least 84% pairwise AA identity). Previously, it was demonstrated that immunity proteins sharing high homology may provide pyocin-silencing functionality of another partner (Dingemans *et al.*, 2016).

To examine to what extent sequence divergence limits the immune function, PmiA homologues from the cluster with *P. fluorescens* Q8r1-96 were selected (*P. brassicacearum* DF41, *P. fluorescens* Pf29Arp, and *Pseudomonas* sp. Pf153), as well as some more distant PmiAs (*Pseudomonas* sp. strains 5 and GM21, *P. fluorescens* AU2390) (Figure 1B). The PmiAs from the Q8r1-96 cluster share 36% to 41% AA identity with PmiA_{Q8r1-96}. This level of sequence conservation stems mainly from the periplasm-exposed regions as well as the transmembrane helices, whereas the amino-terminal signal sequence/transmembrane helix and the cytoplasm-exposed stretches exhibit extensive sequence divergence. By comparison, the ColM domains of the PseuMs of strains DF41, Pf29Arp

and Pf153, display considerably higher sequence conservation (~69% AA identity) with PflM. Pairwise identity with the more distant PmiAs ranges from 18%-31%, corresponding with more diversified ColM domains (35-46% AA identity with PflM).

These six *pmiA* genes were cloned in pJB3Tc20 and introduced in *P. fluorescens* F113, and tested for their capacity to protect from PflM activity, using empty vector and the native *pmiA*_{Q8r196} immunity gene as controls for susceptibility and immunity, respectively (Table 1). Despite only moderate pairwise identities to PmiA_{Q8r1-96}, PmiA_{DF41}, PmiA_{Pf153} and PmiA_{Pf29Arp} provided full cross-immunity to PflM. The more distantly related PmiAs from strains AU2390 and GM21 (with 30.9% and 23.4% AA identity, respectively) could not immunize cells against the bacteriocin, resulting in a similar phenotype as the PmiAs from *P. aeruginosa* NCTC10332 and *P. syringae* DC3000, and the control. Despite barely showing homology with PmiA_{Q8r1-96} (18.4% AA identity), PmiA from *Pseudomonas* sp. 5 conferred an intermediate immunity phenotype (turbid halo).

4 Discussion

In this study we demonstrated that cognate immunity to a large subset of ColM domain-containing *Pseudomonas* bacteriocins is provoked by the poorly conserved transmembrane protein PmiA, unrelated in sequence and topology to the Cmi protein providing immunity to colicin M in *E. coli*. The *pmiA* gene, consistently located downstream of *pseuM* on the same strand, is rarely annotated (Table S3). Previously, immunity to ColM domains in *Burkholderia* burkhocins was equally associated with a transmembrane protein (BmiA) (Ghequire & De Mot, 2015b). However, since ColM domains from *bmiA*- and *pmiA*-associated bacteriocins constitute phylogenetically distant clades, the immunity mechanism from these integral membrane proteins may be different. Further support for the existence of such evolutionarily distinct lipid II-targeting bacteriocin subsets is provided by the observation that a minor group of pseudomonad genes shares a similar immunity gene, as well as a similar ColM-domain-encoding gene, with *Burkholderia*. The gene organization of different ColM domain-hosting bacteriocins and their (putative) immunity genes is summarized in Figure 5.

A conserved Asp residue located in a semi-conserved periplasm-exposed stretch proved not pivotal for the immunity phenotype of PmiA. Possibly, it is required for structural integrity or stability of the immunity protein, and structural analysis can give more insight in this issue. Since the periplasm-exposed moieties of PmiA likely contribute to immunity, it was investigated whether cross-immunity between proteins with different extent of sequence conservation in those regions, may occur. Since this was found to be the case, even between PmiAs sharing less than 40% pairwise amino acid identity, it will be of future interest to determine which residues or structural elements actually contribute to the immunity phenotype. This will render a first indication on how and where the immunity mechanism to this toxin module may be provoked. Previously, attempts to co-crystallize colicin M and Cmi to enlighten how Cmi neutralizes ColM-mediated toxicity proved unsuccessful (Usón *et al.*, 2012).

From an evolutionary point of view, promiscuous protection by immunity genes sharing low homology may have far-reaching consequences towards competitive advantage for producer strains. It may explain why genome analysis in pseudomonads indicated that PseuMs are rather rare, as compared to DNase and RNase bacteriocins (Ghequire & De Mot, 2014). In contrast, immunity to the latter bacteriocins is provided by immunity proteins forming very specifically high-affinity

bacteriocin-immunity complexes, stabilized by conserved hydrogen bonds and hydrophobic interactions (Joshi *et al.*, 2015; Kleanthous *et al.*, 2016; Levin *et al.*, 2009; Li *et al.*, 2004; Meenan *et al.*, 2010). PseuM bacteriocin production may therefore provide limited benefit in most environments, and hence result in lower selection pressure to maintain these bacteriocins in genomes. In addition to this, genome mining also indicated that orphan *pmiA*-like genes occasionally occur in *Pseudomonas* genomes, although it remains to be investigated whether these enable cheating. A similar observation was made for *bmiA* genes in *Burkholderia* (Ghequire & De Mot, 2015b) and S-type pyocin immunity genes (Ghoul *et al.*, 2015).

5 Conflict of Interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

6 Author Contributions

MGKG, RDM conceived and designed the experiments; MGKG and LK performed the experiments; MGKG, LK, and RDM analyzed the data; MGKG and RDM wrote the paper; all authors approved the final version of the manuscript.

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10 Figure and Table legends

Figure 1. Maximum likelihood phylogenetic trees of ColM domains from *pmiA*-associated PseuMs and from related γ -proteobacterial bacteriocins (A), and of associated *Pseudomonas* PmiA immunity proteins (B). Characterized PseuM bacteriocins and cognate PmiAs are colored blue, colicin M (ColM) from *E. coli* is shown in red, and colicin M domain-containing pectocins (PecM1 and PecM2) in pink. PmiAs that were studied for promiscuity are shown in teal. The scale bars represent 0.3 (A) and 0.8 (B) substitutions per site. Bootstrap values (percentages of 1000 replicates) higher than 50 are shown at the branches. (A) For highly homologous colicin M domains (>95% pairwise AA identity), only one representative was included for phylogenetic tree construction. Species abbreviations: Ecoli, *Escherichia coli*; Paer, *Pseudomonas aeruginosa*; Pbra, *Pseudomonas brassicacearum*; Pbre, *Pseudomonas brenneri*; Pcar, *Pectobacterium carotovorum*; Pflu, *Pseudomonas fluorescens*; Pput, *Pseudomonas putida*; Psp, *Pseudomonas* sp.; Psyr, *Pseudomonas syringae*; Psyr cory, *P. syringae* pv. *coryli*; Psyr tom, *P. syringae* pv. *tomato*; Ptol, *Pseudomonas tolaasii*.

Figure 2. Membrane topology model of integral membrane proteins PmiA and BmiA, and membrane-anchored proteins Cmi, PmiB and BmiB, providing immunity to the activity of the ColM domain. PmiA from *Pseudomonas* has four predicted transmembrane helices, the first of which may represent a Sec- or Tat-dependent signal sequence (box). BmiA from *Burkholderia* carries three transmembrane helices. Cmi has a periplasmic YebF domain and is anchored in the inner membrane (IM) via an amino-terminal helix. PmiB and BmiB are equally anchored in the IM, and carry homologous periplasmic domains. “N” depicts the amino-terminus.

Figure 3. Genomic context of orphan *pmiA* genes in pseudomonads. Syntenic regions carrying a *pmiA* homologue associated with a full-length PseuM-encoding gene are shown for *Pseudomonas* sp. strains AU9518 (A), GM55 (B), NFACC04-2 (C), and NFACC51 (D). Synteny between these predicted bacteriocin producers and phylogenetically related cheaters lacking a (functional) bacteriocin gene is highlighted by upstream and downstream ORFs (arrows) shown in the same color. The grey-colored ORFs (D) reflect a local lack of synteny. Remnants of the respective *pseuM* genes are represented by white-coloured rounded shapes corresponding to a frameshifted *pseuM* (position marked by a dotted vertical line) or a truncated residual N- or C-terminal fragment. For *Pseudomonas* strains ERGS4:02 (E) and Leaf48 (F), no strain with a closely related *pmiA* homologue and syntenic flanking region could be identified. Species abbreviations: Pfre, *Pseudomonas frederikbergensis*; Plun, *Pseudomonas lundensis*; Pwei, *Pseudomonas weihenstephanensis*. Other abbreviations are as in Figure 1.

Figure 4. Multiple sequence alignment of the semi-conserved stretch in PmiA proteins, predicted to be oriented towards the periplasmic space. *Pseudomonas* species abbreviations are as in Figure 1. Grey shading reflects the degree of conservation. The conserved Asp residue that was mutated is boxed in green.

Figure 5. Schematic gene organizations of ColM domain-containing bacteriocins and (putative) immunity genes (if present). The arrows correspond with the gene orientations and the color legend describes the indicated domains and type of immunity gene. Predicted immunity genes, based on homology with characterized immunity factors, are delineated by a dashed line. Species abbreviations are as in Figure 1. Other abbreviations: Bamb, *Burkholderia ambifaria*; Bglu, *Burkholderia glumae*.

Table 1. Heterologous expression of *pmiA* genes in PseuM-susceptible strains. Indicator strains were transformed with shuttle vector pJB3Tc20 (control, without insert) or with pJB3Tc20-derived plasmids equipped with different *pmiA* genes, and tested for (altered) susceptibility to their respective PseuM via bacteriocin spot assay. +, sensitive; -, insensitive; T, turbid halo. The species used as source of immunity protein are specified with the respective strain names indicated as PmiA subscript (NA, not applicable). Asterisks define mutated immunity proteins (Asp to Ala). Strain LMG 5456 carrying a plasmid equipped with *pmiA*_{Q8r1-96} could not be obtained.

Indicator strain	PmiA immunity protein		PseuM bacteriocin		
	Species	Gene product	PaeM <i>P. aeruginosa</i> NCTC10332	PflM <i>P. fluorescens</i> Q8r1-96	PsyM <i>P. syringae</i> pv. <i>tomato</i> DC3000
<i>P. aeruginosa</i> CPHL12447	<i>P. aeruginosa</i>	PmiA _{NCTC10332}	-		
		PmiA* _{NCTC10332}	-		
	<i>P. fluorescens</i>	PmiA _{Q8r1-96}	+		
	<i>P. syringae</i>	PmiA _{DC3000}	+		
	NA	Control	+		
<i>P. fluorescens</i> F113	<i>P. aeruginosa</i>	PmiA _{NCTC10332}		+	
	<i>P. brassicacearum</i>	PmiA _{DF41}		-	
	<i>P. fluorescens</i>	PmiA _{Q8r1-96}		-	
		PmiA* _{Q8r1-96}		-	
		PmiA _{AU2390}		+	
		PmiA _{Pf29Arp}		-	
	<i>P. syringae</i>	PmiA _{DC3000}		+	
	<i>Pseudomonas</i> sp.	PmiA ₅		T	
		PmiA _{GM21}		+	
		PmiA _{Pf153}		-	
	NA	Control		+	
<i>P. syringae</i> LMG 5456	<i>P. aeruginosa</i>	PmiA _{NCTC10332}			+
	<i>P. syringae</i>	PmiA _{DC3000}			-
		PmiA* _{DC3000}			-
	NA	Control			+

11 Supplementary tables and figures

Table S1. Strains and plasmids used in this study.

Table S2. Primers used in this study.

Table S3. Colicin M-like bacteriocin genes present in genome sequences of pseudomonads. NA: not annotated. PseuM genes present in gene context consisting of phage-like genes are marked with “Y” (phage-like and tailocin context), others with “N”. Pseudomonads containing a putative orphan *pmiA* are colored red.

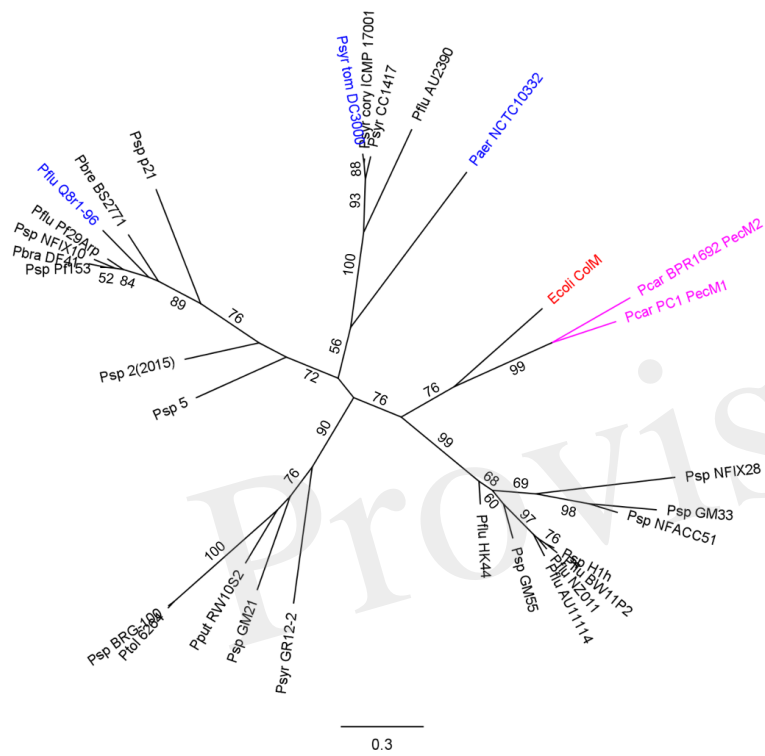
Figure S1. (A) Multiple sequence alignment of ColM domains derived from unique PseuM proteins from *Pseudomonas* spp. Abbreviations for species names are as in Figure 1. Other abbreviations: Pced, *Pseudomonas cedrina*; Pcor ory, *Pseudomonas coronafaciens* pv. *oryzae*; Pden, *Pseudomonas denitrificans*; Pkor, *Pseudomonas koreensis*; Ppro, *Pseudomonas protegens*; Psyr ace-cori-mors-syr-the, *Pseudomonas syringae aceris-coriandricola-morsprunorum-syringae-theae*; Pthi, *Pseudomonas thivervalensis*; Pver, *Pseudomonas veronii*. Grey shading reflects the degree of conservation. (B) Unrooted maximum likelihood phylogenetic tree of ColM domains of pseudomonads (black) from panel (A) and previously characterized ColM domain-carrying bacteriocins from other β - and γ -proteobacteria: colicin M (ColM, red) from *E. coli*, pectocin M1 and M2 from *Pectobacterium* spp. (PecM1/PecM2, pink), burkhocin M1 and M2 from *Burkholderia* spp. (BurM1/BurM2, grey). *Pseudomonas* abbreviations as in (A). The clade of *Pseudomonas* ColM domains from bacteriocins linked with a *pmiA*-type immunity gene and *pmiB*-type immunity gene are in a red arc and a blue ellipse, respectively. Other abbreviations: Bamb, *Burkholderia ambifaria*; Ecoli, *Escherichia coli*; Pcar, *Pectobacterium carotovorum*. The scale represents 0.6 substitutions per site. Bootstrap values (percentages of 1000 replicates) are not shown for visibility.

Figure S2. Multiple sequence alignment of BmiB (*Burkholderia* spp.) and unique PmiB (*Pseudomonas* spp.) proteins. Grey shading indicates the degree of conservation. Abbreviations for *Burkholderia* species names: Bubo, *Burkholderia ubonensis*; Bcon, *Burkholderia contaminans*; Bamb, *Burkholderia ambifaria*; Bokl, *Burkholderia oklahomensis*; Bcep, *Burkholderia cepacia*; Bcen, *Burkholderia cenocepacia*; Bpyr, *Burkholderia pyrrocinia*; Burk, *Burkholderia* sp. *Pseudomonas* abbreviations are as in Figure 1 and Figure S1. Predicted transmembrane helices are boxed in blue, Sec- or Tat-dependent signal sequences in red and lipoprotein signal sequences in yellow. Two conserved cysteine residues are highlighted in green.

Figure S3. Multiple sequence alignment of unique PmiA proteins. Predicted transmembrane helices are boxed in blue, Sec- or Tat-dependent signal sequences in red. Grey shading indicates the degree of conservation. *Pseudomonas* species abbreviations are as in Figure S1. This multiple sequence alignment was used for construction of phylogenetic tree in Figure 1B.

Figure 01.TIF

A



B

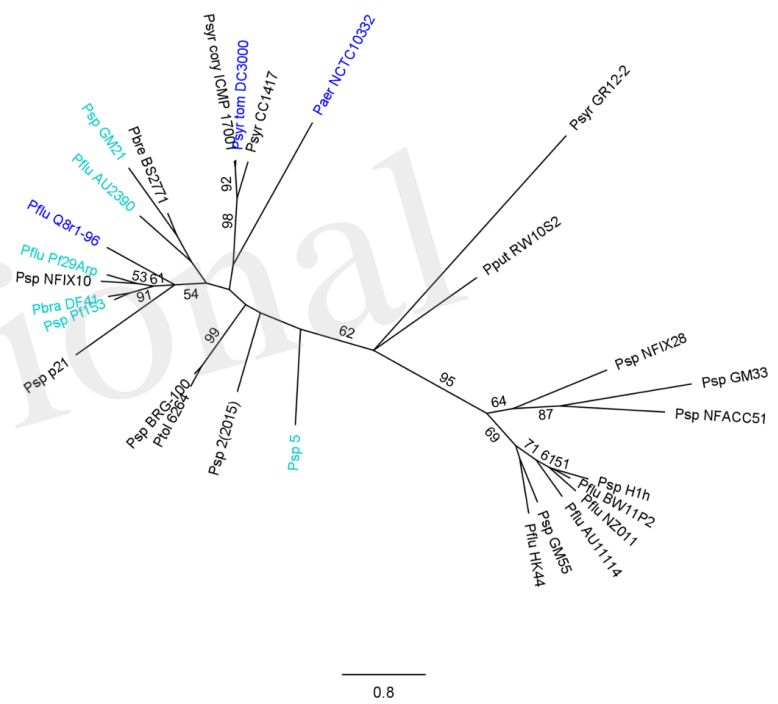


Figure 02.TIF

Provisional

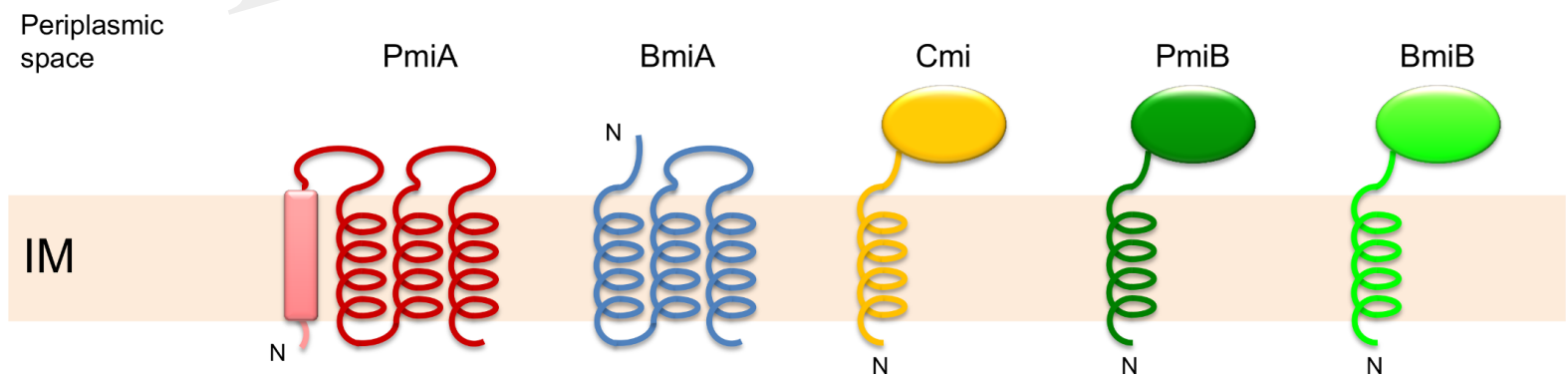


Figure 03.TIF

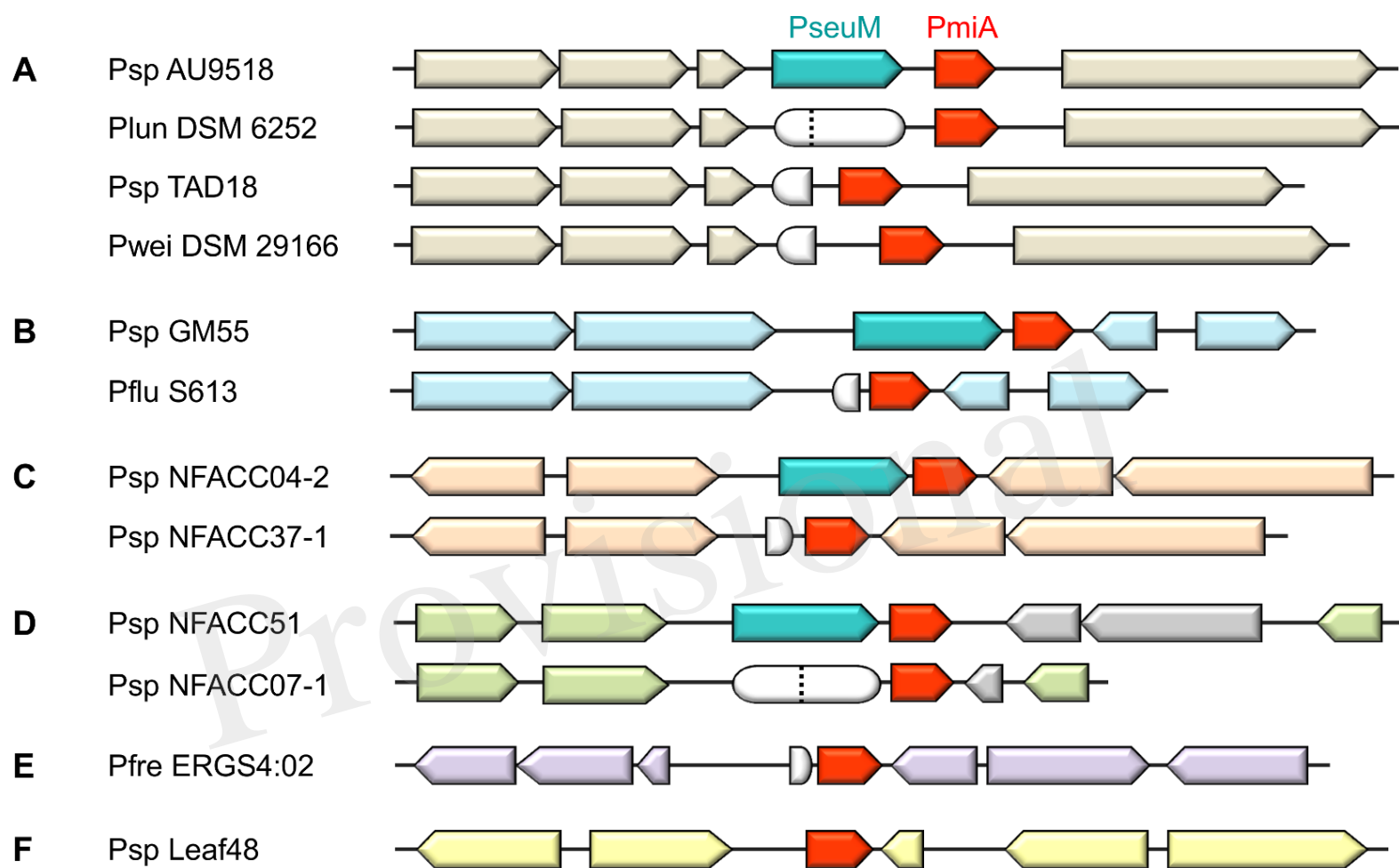


Figure 04.TIF

Psp BRG-100	L E N P N D P S -	D T V G V P
Ptol 6264	L E N P N D P S -	D T A G V P
Pput RW10S2	Q G D I D L G G -	E N S G - P
Psp GM21	P P N S K D L S -	D T S G L S
Psyr GR12-2	P V I A H S G R	D E I G D D
Psp GM33	K N S I Y P T M Q	D T I A I P
Psp NFIX28	S S G K Y S S E A	D S L A I P
Psp NFACC51	E S D L Y A V E H	D T I A I P
Psp H1h	R A E I Y D V N Q	D S I G L P
Pflu BW11P2	E S G I Y D V N Q	D S I G L P
Pflu NZ011	E A Q I Y D A K Q	D S I G I P
Pflu AU11114	E L K K Y Q M D R	D S V G L P
Psp GM55	S A G I Y D I D R	D S I G I P
Pflu HK44	S A G V Y P I H Q	D S I G L P
Psp 2(2015)	P E N L D D P L -	D T R G V P
Psp 5	P P N P D D P Q -	D T Y G L P
Psp Pf153	T E N P K N L S -	D T Q G I P
Pbra DF41	S E N P N D L S -	D T R G I P
Psp NFIX10	V E N P R D L N -	D T Q G I P
Pflu Pf29Arp	V E N P G N P N -	D T Y G L P
Pflu Q8r1-96	P E N P G N P S -	D T S G L P
Pbre BS2771	P E N P A D L S -	D T S G L P
Psp p21	L E N P G D P H -	D T S G I P
Pflu AU2390	A E N P S N L S -	D T T G V P
Paer NCTC10332	S E G N V N P D -	D T T G V P
Psyr CC1417	A E N P L S P S -	D T Q G L P
Psyr cory ICMP 17001	S E N P L Q P S -	D T Y G L P
Psyr tom DC3000	S E N P L Q P S -	D T Y G L P

Figure 05.TIF

Provisional

